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<p>2003-610325/58</p> <p>RHODIANYL SNC</p> <p>2001.12.17 2001-016322(+2001FR-016322) (2003.06.20) C08L 77/06, C08G 69/06, C08J 5/18 (C08L 77/06, 77.00)</p> <p>Thermoplastic polymer composition, used e.g. for production of moulded products, film or fibres, contains a functionalised, hyperbranched copolyamide as rheology modifier</p> <p>C2003-166585</p>	<p>A23 F01</p> <p>RHOD 2001.12.17</p> <p>*FR 2833603-A1</p> <p>NOVELTY</p> <p>Functionalised, hyper-branched copolyamides are used as rheology modifiers in thermoplastic polymer compositions.</p>	<p>DETAILED DESCRIPTION</p> <p>A thermoplastic polymer composition (TPC) comprising a thermoplastic polymer-based matrix (M) and rheology-modifying additive(s) (PAH) consisting of functionalised, hyperbranched copolyamide(s) obtained by reaction of</p> <ul style="list-style-type: none"> (a) monomer(s) of formula A-R-B_f (I), possibly (b) difunctional spacer monomer(s) of formula A'-R'-B' (II) or the corresponding lactams, possibly (c) core monomer(s) of formula R¹(B)_n (III) and (d) chain-regulating monomer(s) of formula R²-A (IV), at least 50% of the terminal groups in the copolyamide being functionalised by R². 	<p>USE</p> <p>Functionalised hyperbranched copolyamides (as described) are used as rheology modifiers for thermoplastic polymer matrices (claimed). Polymers modified with these copolyamides are used e.g. for the production of moulded articles, yarn, fibres, film and filaments.</p>
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ADVANTAGE

Functionalised hyperbranched copolyamides enable the controlled modification of the rheological properties of thermoplastics (especially melt viscosity) without adverse effects on mechanical properties such as impact strength and without changing the chemical structure of the matrix or reducing its mol. wt. These copolyamides are also readily dispersible in the matrix polymer.

EXAMPLE

A preheated (70°C) reactor was charged in succession with 1867.4 g hexadecylamine (melt; purity 90%), 525.5 g ε-caprolactam, 840.6 g 5-amino-isophthalic acid, 162.5 g benzene-1,3,5-tricarboxylic acid and 6.1 g 50% aqueous hypophosphorous acid, then the mixture was stirred at 260°C for 30 minutes and vacuum was applied progressively over 1 hour, to give 2946 g hyperbranched copolyamide (PAHB-C16/2) and 155.2 g distillate. The product showed mol. wts. of 4890, 7750 and 11440 (Mn, Mw, Mz), a mol. wt. ratio of 1.58, an amine end group content of 3.1 meq/kg and an acid end group content of 7.7 meq/kg. A mixture of polyamide 6/50% glass fibres and 2 wt% PAHB-C16/2 showed a melt viscosity (flow spiral; 300°C) of 430 (339) mm and a

moisture content (before moulding) of 0.26 (0.20) %. Injection moulded test samples showed a mol. wt. (PA 6,6) of 63550 (65250), an impact strength (un-notched) of 76.3 (82.4) kJ/m², an impact strength (notched) of 10.3 (11.0) kJ/m², a flexural modulus of 11500 (11800) N/mm², a flexural strength of 262 (280) N/mm² and a heat distortion point of 245 (248) °C. Values in brackets are for unmodified PA66/50% glass fibres; the change in mol. wt. caused by the modifier was -2.6%. Other combinations of melt viscosity and other properties were obtained by using different types and amounts of copolyamide modifier.

DEFINITIONS

Preferred Definitions:
R, R' = linear or branched aliphatic groups and/or cycloaliphatic groups and/or mono- or poly-nuclear aromatic groups and/or aryl-aliphatic groups (all optionally substituted and/or containing hetero-atoms), with aromatic groups especially preferred for R in formula (I);
A, A' = amino, amine salt, acid, ester, acid halide or amide groups, preferably carboxylic or amino functions, with amino

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especially preferred for A in (I);
B, B' = acid, ester, acid halide, amide, amino or amine salt groups,
preferably carboxylic or amino functions, with carboxylic
especially preferred for B in (I);
 $f = 2$

TECHNOLOGY FOCUS

Polymer - Preferred Compositions: Compositions containing no

PAHB additives which cause a reduction of 7% or more in the mol.
wt. of the matrix (M) (using a specified method for mol. wt.

determination, preferably carried out on the composition to be
analysed and the reference composition, both extruded, solidified and
granulated and possibly subjected to a quantitative fluidity test (Qf)).

Preferred compositions: The composition contains 0.1-50 (preferably
1-20, especially 2-10) wt% PAHB. Other components include
reinforcing materials and/or fillers such as glass, aramid, ceramic or
carbon fibres, mineral fillers, fillers based on thermosets and
powdered fillers such as talcum. Preferred Copolyamides:

Hyperbranched copolyamides (PAHB) with an acid or amine end-
group content of 0-25 meq/kg, containing monomers (III) with a mol

ratio of (III)/(I+II+IV) = not more than 1/150, preferably not more
than 1/100, especially not more than 1/50.

Preferred Matrix Polymers: Twenty-two polymer types listed,
including e.g. polyolefins, polyesters, polyalkylene oxides,
polyphenylene, polyvinyl acetates, polyvinyl halides, polyamides,
polycarbonates, acrylic polymers, cellulose or its derivatives and
thermoplastic copolymers.

Preferred polymers are (co)polyamides such as polyamide (PA) 6, 6.6,
4, 11, 12, 4-6, 6-10, 6-12, 6-36 and/or 12-12, especially PA 6 with a
relative viscosity (1 % in sulfuric acid at 25°C) of more than 3.5,
preferably more than 3.8, or PA 6.6.

Preferred Articles: Moulded products, yarn, fibres, film or filaments
obtained from TPC.

Organic Chemistry - Preferred Monomers: Monomer (I) comprises 5-
amino-isophthalic, 6-amino-undecanoic, 3-aminopimelic, aspartic, 3,5-
diaminobenzoic and/or 3,4-diaminobenzoic acid;
(II) comprises ϵ -caprolactam and/or aminocaproic acid, p- or m-
aminobenzoic acid, 11-amino-undecanoic acid, lauryl-lactam and/or
the corresponding amino-acid and/or 12-aminododecanoic acid;

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(III) comprises 1,3,5-benzenetricarboxylic acid, 2,2,6,6-tetra-(β -carboxyethyl)-cyclohexanone, 2,4,6-tri-(aminocaproic acid)-1,3,5-triazine and/or 4-aminoethyl-octane-1,8-diamine;
(IV) comprises n-hexadecylamine, n-octadecylamine, n-dodecylamine and/or benzylamine.
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